

**K240901 Stethophone**Sep 19, 2024  
170 days to decisionK240901 · Product code: **DQD** · CardiovascularSource: <https://www.510kdatabase.net/k240901/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stethoscope, Electronic (DQD)
Date received	Apr 2, 2024
Decision date	Sep 19, 2024
Days to decision	170 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Sparrow Acoustics, Inc.</b>
Location	Lucasville, CA
Contact	Nadia Ivanova
510(k) history	4 submissions · 4 cleared · 2023-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k240901/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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